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=> s benzenesulfonylcarbomoyl and (erythromycin or azithromycin or homoerythromycin)
0 BENZENESULFONYLCARBOMOYL

18623 ERYTHROMYCIN

3796 AZITHROMYCIN

99 HOMOERYTHROMYCIN

L1 0 BENZENESULFONYLCARBOMOYL AND (ERYTHROMYCIN OR AZITHROMYCIN) OR HOMOERYTHROMYCIN)

=> s benzenesulfonyl and carbamoyl

9083 BENZENESULFONYL

24918 CARBAMOYL

L2 291 BENZENESULFONYL AND CARBAMOYL

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=> s 12 and (erythromycin or azithromycin or homoerythromycin)
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             99 HOMOERYTHROMYCIN
L3
              2 L2 AND (ERYTHROMYCIN OR AZITHROMYCIN OR HOMOERYTHROMYCIN)
=> d 13 1-2
     ANSWER 1 OF 2 CA COPYRIGHT 2008 ACS on STN
1.3
AN
     140:407071 CA
     Substituted 9a-N-[N'-(benzenesulfonyl)carbamoyl
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     -y-aminopropyl] and 9a-N-[N'-(b-cyanoethyl)-N'-(benzenesulfonyl)
     carbamoyl-y-aminopropyl] derivatives of 9-de-oxo-9-dihydro-9a-aza-
     9a-homo-erythromycin A and 5-0-desosaminyl-9-de-oxo-9-dihydro-9a-
     aza-homo-erythronolide A
     Kujundzic, Nedjeljko; Bukvic Krajacic, Mirjana; Brajsa, Karmen
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     Pliva D.D., Croatia
so
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DT
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LΑ
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              THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L3
     ANSWER 2 OF 2 CA COPYRIGHT 2008 ACS on STN
AN
     137:201527 CA
     Preparation of 9a-N-[N'-(phenylsulfonyl)carbamoyl] derivatives
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     of 9-deoxo-9-dihydro-9a-aza-9a-homoerythromycin A and of
     5-0-desosaminyl-9-deoxo-9-dihydro-9a-aza-9a-homoerythronolide A as
     antibacterial agents
IN
     Kujundzic, Nedjeljko; Bukvic Krajacic, Mirjana; Dumic, Miljenko;
     Hasenohrl, Andrea
PΑ
     Pliva D.D., Croatia
SO
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
DT
     Patent
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    CASREACT 137:201527; MARPAT 137:201527
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=> d l3 1-2 an ab

L3 ANSWER 1 OF 2 CA COPYRIGHT 2008 ACS on STN

AN 140:407071 CA

AΒ The invention relates to substituted 9a-N-[N'-(benzenesulfonyl) carbamoyl- γ -aminopropyl] and 9a-N-[N'-(β -cyanoethyl)-N'-(benzenesulfonyl) carbamoyl- γ -aminopropyl] derivs. of 9-de-oxo-9-dihydro-9a-aza-9a-homo-erythromycin A and 5-0-desosaminyl-9-de-oxo-9-dihydro-9a-aza-9a-homo-erythronolide A, novel semisynthetic macrolide antibiotics of the azide series, of the formula I, wherein R represents H or cladinosyl moiety, R1 represents H or β -cyanoethyl group and R2 represents H or fluoro, chloro and Me group, and pharmaceutically acceptable salts thereof with inorg. or organic acids, to the process for the preparation of pharmaceutical compns. as well as to the use their compns. for sterilization rooms and medical instruments as well as for protection of wall and wooden coatings. Test substances of title macrolide glycosides were active on susceptible strains of S. pyogenes (MIC 0.125 to 4.0 mg/L), and on susceptible strains on S. pneumoniae (MIC 0.125 to 8.0 mg/L). MIC values on susceptible S. aureus strains were from 1 to 16 mg/L. Substances showed strong antimicrobial activities on most tested Gram neg. strains; M. catarrhalis MIC from 0.25 to 16 mg/L, E. coli from 8 to 16 mg/L, E. faecalis from 2 to 8 mg/L.

L3 ANSWER 2 OF 2 CA COPYRIGHT 2008 ACS on STN

AN 137:201527 CA

AB The invention relates to 9a-N[N'-(phenylsulfonyl)carbamoyl]
derivs. of 9-deoxo-9-dihydro-9a-aza-homoerythromycin A and
5-0-desosaminyl-9-deoxo-9-dihydro-9a-aza-9a-homoerythronolide A, novel
semisynthetic macrolide antibiotics from the class of azalides, of the
general formula I wherein R1 denotes H, C1-C4alkyl or halogen and R2
denotes H or cladinosyl radical, to their pharmaceutically acceptable
addition salts with inorg. or organic acids, to intermediates and methods for
their preparation, to a process for the preparation of pharmaceutical compns.

well as to the use of pharmaceutical compns. in the treatment of bacterial infections. Thus, 9-deoxo-9-dihydro-9a-N-[N'-(4-chlorobenzenesulfonyl) carbamoyl]-9a-aza-9a-homoerythromycin A was prepared by

as

condensation reaction of 9-deoxo-9-dihydro-9a-azahomoerythomycin A with 4-chlorobenzensulfonyl isocyanate. These glycoside macrolides were used as antibacterial agents (no data).